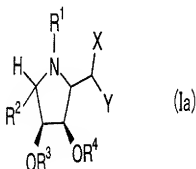


### Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

#### Listing of Claims:

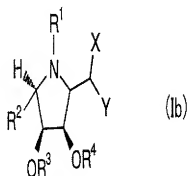
1. (Canceled)
2. (Currently Amended) ~~The A compound according to claim 1 or a salt thereof,~~  
~~wherein the configuration of the formula (I) is represented by the following formula (Ia)~~  
or a salt thereof:



wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$ ,  $X$ , and  $Y$  are as defined in claim 1

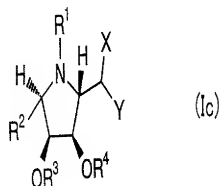
wherein  $R^1$  represents a hydrogen atom, a  $C_{1-10}$  alkyl group optionally having a substituent, or a protecting group of N;  $R^2$  represents a  $C_{1-10}$  alkyl group optionally having a substituent or a  $C_{2-10}$  alkenyl group optionally having a substituent;  $R^3$  and  $R^4$  independently represent a hydrogen atom or a protecting group of hydroxyl group;  $X$  represents  $-N(R^5)R^6$  or a residue of amino acid or of an amino group of a peptide wherein  $R^5$  and  $R^6$  independently represent a hydrogen atom, a  $C_{1-10}$  alkyl group optionally having a substituent, or a  $C_{3-12}$  cycloalkyl group optionally having a substituent; and  $Y$  represents a hydrogen atom,  $-CH_2NH_2$ ,  $-CONH_2$ , or  $-COOH$ .

3. (Currently Amended) The compound according to claim  $[[1]]$  2 or a salt thereof, wherein the configuration of the formula (†) (Ia) is represented by the following formula (Ib):



wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$ , X, and Y are as defined in claim  $[[1]]$  2.

4. (Currently Amended) The compound according to claim  $[[1]]$  2 or a salt thereof, wherein the configuration of the formula (†) (Ia) is represented by the following formula (Ic):



wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$ , X, and Y are as defined in claim  $[[1]]$  2.

5. (Currently Amended) The compound according to claim  $[[1]]$  2 or a salt thereof, wherein  $R^2$  represents  $-CH_2OR^{12}$  wherein  $R^{12}$  represents a hydrogen atom or a protecting group of hydroxyl group.

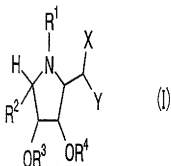
6. (Currently Amended) An inhibitor composition of sugar chain related enzymes which comprises, as an active ingredient, the compound of claim  $[[1]]$  2 or a salt thereof.

7. (Currently Amended) A ~~medicine~~ pharmaceutical composition which comprises, as an active ingredient, the compound of claim  $[[1]]$  2 or a salt thereof.

8. (Currently Amended) The ~~medicine~~ pharmaceutical composition according to claim 7 which is a medicine for ~~the~~ therapy or prevention of diseases associated with sugar chain related enzymes.

9. (Currently Amended) The ~~medicine~~ pharmaceutical composition according to claim 7 which ~~is used as~~ is an antiviral agent, an anticancer agent, or an immunostimulant agent.

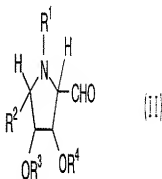
10. (Currently Amended) A method for producing a compound represented by the formula (I);



wherein R<sup>1</sup> represents a hydrogen atom, a C<sub>1-10</sub> alkyl group optionally having a substituent, or a protecting group of N; R<sup>2</sup> represents a C<sub>1-10</sub> alkyl group optionally having a substituent or a C<sub>2-10</sub> alkenyl group optionally having a substituent; R<sup>3</sup> and R<sup>4</sup> independently represent a hydrogen atom or a protecting group of hydroxyl group; X represents -N(R<sup>5</sup>)R<sup>6</sup> or a residue of amino acid or of an amino group of a peptide wherein

R<sup>5</sup> and R<sup>6</sup> independently represent a hydrogen atom, a C<sub>1-10</sub> alkyl group optionally having a substituent, or a C<sub>3-12</sub> cycloalkyl group optionally having a substituent; and Y represents a hydrogen atom, -CH<sub>2</sub>NH<sub>2</sub>, -CONH<sub>2</sub>, or -COOH;

according to claim 1 which comprises a step of comprising reacting a compound represented by the formula (II):

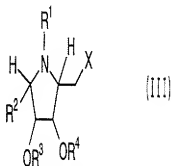


wherein R<sup>1</sup> represents a hydrogen atom, a C<sub>1-10</sub> alkyl group optionally having a substituent, or a protecting group of N; R<sup>2</sup> represents a C<sub>1-10</sub> alkyl group optionally having a substituent or a C<sub>2-10</sub> alkenyl optionally having a substituent; and R<sup>3</sup> and R<sup>4</sup> independently represent a hydrogen atom or a protecting group of hydroxyl group;

with a compound represented by the formula X-H:

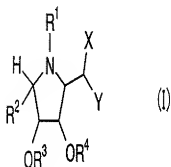
wherein X represents -N(R<sup>5</sup>)R<sup>6</sup> or a residue of amino acid or of an amino group of a peptide, and R<sup>5</sup> and R<sup>6</sup> independently represent a hydrogen atom, a C<sub>1-10</sub> alkyl group optionally having a substituent, or a C<sub>3-12</sub> cycloalkyl group optionally having a substituent;

in the presence of a reducing agent, to produce a compound represented by the formula (III):



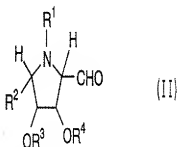
wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup> and X are as defined above.

11. (Currently Amended) A method for producing a compound represented by the formula (I):



wherein R<sup>1</sup> represents a hydrogen atom, a C<sub>1-10</sub> alkyl group optionally having a substituent, or a protecting group of N; R<sup>2</sup> represents a C<sub>1-10</sub> alkyl group optionally having a substituent or a C<sub>2-10</sub> alkenyl group optionally having a substituent; R<sup>3</sup> and R<sup>4</sup> independently represent a hydrogen atom or a protecting group of hydroxyl group; X represents -N(R<sup>5</sup>)R<sup>6</sup> or a residue of amino acid or of an amino group of a peptide wherein R<sup>5</sup> and R<sup>6</sup> independently represent a hydrogen atom, a C<sub>1-10</sub> alkyl group optionally having a substituent, or a C<sub>3-12</sub> cycloalkyl group optionally having a substituent; and Y represents a hydrogen atom, -CH<sub>2</sub>NH<sub>2</sub>, -CONH<sub>2</sub>, or -COOH;

according to claim 1 which comprises a step of comprising reacting a compound represented by the formula (II):

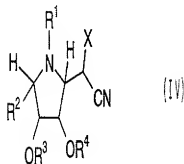


wherein  $R^1$  represents a hydrogen atom, a  $C_{1-10}$  alkyl group optionally having a substituent, or a protecting group of N;  $R^2$  represents a  $C_{1-10}$  alkyl group optionally having a substituent or a  $C_{2-10}$  alkenyl optionally having a substituent; and  $R^3$  and  $R^4$  independently represent a hydrogen atom or a protecting group of hydroxyl group;

with a compound represented by the formula X-H:

wherein X represents  $-N(R^5)R^6$  or a residue of amino acid or of an amino group of a peptide, and  $R^5$  and  $R^6$  independently represent a hydrogen atom, a  $C_{1-10}$  alkyl group optionally having a substituent, or a  $C_{3-12}$  cycloalkyl group optionally having a substituent;

and a cyanation agent in the presence of Lewis acid, to produce a compound represented by the formula (IV):



wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$  and X are as defined above, and converting the nitrile group to an amino group or amide group.

12. (New) The compound according to claim 3 or a salt thereof, wherein  $R^2$  represents  $-CH_2OR^{12}$  wherein  $R^{12}$  represents a hydrogen atom or a protecting group of hydroxyl group.

13. (New) The compound according to claim 4 or a salt thereof, wherein  $R^2$  represents  $-CH_2OR^{12}$  wherein  $R^{12}$  represents a hydrogen atom or a protecting group of hydroxyl group.

14. (New) A method of treating diseases associated with sugar chain related enzymes comprising administering a therapeutically effective amount of the compound according to claim 2 or a salt thereof to a mammal.

15. (New) A method of treating diseases associated with sugar chain related enzymes comprising administering a therapeutically effective amount of the compound according to claim 3 or a salt thereof to a mammal.

16. (New) A method of treating diseases associated with sugar chain related enzymes comprising administering a therapeutically effective amount of the compound according to claim 4 or a salt thereof to a mammal.

17. (New) A method of administering an antiviral agent, an anticancer agent, or an immunostimulant agent to a mammal comprising administering a compound according to claim 2 or a salt thereof to the mammal.

18. (New) A method of administering an antiviral agent, an anticancer agent, or an immunostimulant agent to a mammal comprising administering a compound according to claim 3 or a salt thereof to the mammal.

19. (New) A method of administering an antiviral agent, an anticancer agent, or an immunostimulant agent to a mammal comprising administering a compound according to claim 4 or a salt thereof to the mammal.

20. (New) The method according to claim 14 wherein the mammal is a human.

21. (New) The method according to claim 15 wherein the mammal is a human.